

A Nonpeptidic Cathepsin S Activity-Based Probe for Noninvasive Optical Imaging of Tumor-Associated Macrophages

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SUMMARY

Macrophage infiltration into tumors has been correlated with poor clinical outcome in multiple cancer types. Therefore, tools to image tumor-associated macrophages could be valuable for diagnosis and prognosis of cancer. Herein, we describe the synthesis and characterization of a cathepsin S-directed, quenched activity-based probe (qABP), BMV083. This probe makes use of an optimized non-peptidic scaffold leading to enhanced *in vivo* properties relative to previously reported peptide-based probes. In a syngeneic breast cancer model, BMV083 provides high tumor-specific fluorescence that can be visualized using noninvasive optical imaging methods. Furthermore, analysis of probe-labeled cells demonstrates that the probe primarily targets macrophages with an M2 phenotype. Thus, BMV083 is a potential valuable *in vivo* reporter for tumor-associated macrophages that could greatly facilitate the future studies of macrophage function in the process of tumorigenesis.

INTRODUCTION

The cysteine cathepsin family contains 11 members that play important roles in protein breakdown in the lysosome (Turk et al., 2001). Several family members have also been shown to play functional roles in the onset and progression of diverse diseases, ranging from Alzheimer's disease to cancer (Conus and Simon, 2010; Mohamed and Sloane, 2006; Mueller-Steiner et al., 2006; Reiser et al., 2010). Because the activity of these proteases is highly regulated and dependent on posttranslational maturation of the proenzyme, tools that can report on their activity levels have been essential to understanding their biolog-

ical function in disease pathology. In particular, a number of activity-based probes (ABPs) have been developed that allow the direct profiling of cysteine cathepsin activity levels *in vivo*. Recently, we developed a near-infrared fluorophore-labeled quenched ABP (NIRF qABP), GB137, containing a peptide recognition scaffold linked to an acyloxymethyl ketone (AOMK) electrophile (Blum et al., 2007). This intrinsically nonfluorescent probe produces a fluorescent signal upon nucleophilic displacement of the quencher group by the active site cysteine of the proteolytically active cysteine cathepsin (Figure 1A). Because high levels of cysteine cathepsin activity are associated with many tumor types, the probe can be used for direct noninvasive optical imaging applications. Furthermore, because the probes are activity dependent, they can also be used to noninvasively measure the efficacy of small-molecule therapeutics. In contrast to fluorogenic cathepsin substrates that have also been successfully used for live animal imaging experiments (Gounaris et al., 2008; Mahmood and Weissleder, 2003), with the most recent addition being a cathepsin S-selective substrate (Caglic et al., 2011), the irreversible nature of the ABPs allows subsequent downstream identification of the targeted protease and correlation with histological and cell biological assessments of the *in vivo* tumor microenvironment.

The primary *in vivo* targets of our first-generation qABP were cathepsin B, S, and L (Blum et al., 2005; Blum et al., 2007). Although important roles in tumor development have been described for all three of these cysteine cathepsins (Gocheva and Joyce, 2007), cathepsins B and L, like most members of the cysteine cathepsin family, are ubiquitously expressed (Conus and Simon, 2010). Cathepsin S, however, is most abundantly expressed in antigen-presenting cells (APCs), where it plays a major role in MHC II antigen presentation (Zavaski-Bergant and Turk, 2006). Macrophages are professional APCs and are, therefore, key players in immunity. They have a variety of functions depending on their activation state—classically activated (M1) or alternatively activated (M2). Macrophages can also be classified into three groups according to their homeostatic functions: host defense (classically activated

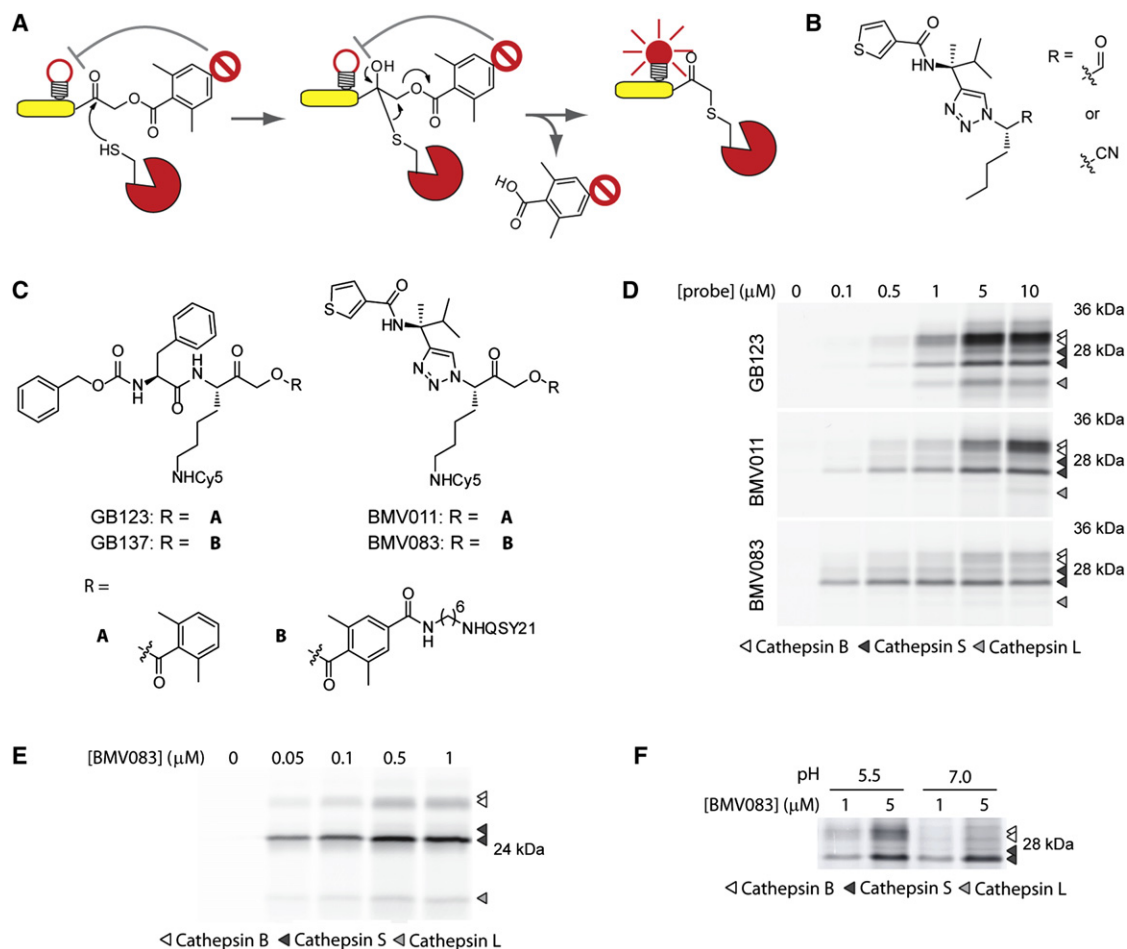


Figure 1. Nonpeptidic Cysteine Cathepsin Activity-Based Probes

(A) Schematic presentation of the mechanism of action of a quenched ABP.

(B) Structure of the cathepsin S-selective aldehyde and nitrile inhibitors reported by the Ellman laboratory.

(C) Structures of the peptidic activity-based probes GB123 and the quenched GB137 and the nonpeptidic probes BMV011 and the quenched BMV083.

(D) Labeling profile of GB123, BMV011, and BMV083 in living RAW cells. Cells were exposed to the indicated concentrations of probe for 3 hr, before being harvested, washed, and lysed. Total protein (40 μg) was resolved on 15% SDS-PAGE, and fluorescently labeled proteins were visualized by in-gel fluorescence scanning.

(E) Labeling profile of BMV083 in living human primary macrophages. Cells were exposed to the indicated concentrations of BMV083 for 3 hr, before being harvested, washed, and lysed. Total protein (40 μg) was analyzed as described above.

(F) BMV083 labeling of RAW cell lysate (35 μg total protein) at pH 5.5 and 7.0 with indicated concentrations of probe for 1 hr. Labeled proteins were analyzed as described above.

See also Figures S1, S2, S3, and S4.

macrophages), wound healing (wound-healing macrophages), and immune regulation (regulatory macrophages) (Mosser and Edwards, 2008). However, macrophages display a high degree of plasticity, and activation states can change in response to stimuli from their environment. Furthermore, macrophages can have a blend of characteristics of multiple groups. One such type of macrophage is the tumor-associated macrophage (TAM), which displays characteristics of both wound-healing and regulatory macrophages and plays important roles in tumorigenesis by promoting angiogenesis, tumor growth, and invasiveness. These macrophages are recruited to the tumor site and are stimulated by factors in the tumor microenvironment, including the cytokine IL-4, which induces cysteine cathepsin activity (Gocheva et al., 2010). In human studies, TAM infiltration

in tumors has been associated with poor prognosis, for example in high-risk breast cancers (Mukhtar et al., 2011).

The development of imaging tools to identify TAM infiltration in tumors could lead to clinical applications for the treatment and prognosis of cancer. Because of its confined expression, probes that are designed to target cathepsin S are likely to provide improved contrast for areas with stimulated macrophages compared to more broad-spectrum probes that also target other cysteine cathepsins that have a broader expression profile. Herein, we describe the synthesis and characterization of a cathepsin S-directed, nonpeptidic NIRF qABP with improved in vivo properties relative to previously reported peptide-based probes. We use this optimized cathepsin S probe for noninvasive optical imaging of a syngeneic mouse model of breast cancer.

Fluorescence-activated cell sorting (FACS) experiments identified specific subsets of myeloid-derived cells with an M2 macrophage phenotype as the *in vivo* cellular source of cysteine cathepsin activity responsible for probe fluorescence. These data demonstrate the potential value of our ABP for the classification of tumor-associated myeloid derived cells based on *in vivo* protease activity.

RESULTS AND DISCUSSION

Synthesis and In Vitro Characterization of BMV083

We set out to develop ABPs that would have a high degree of selectivity for cathepsin S that could be used for noninvasive imaging applications. Furthermore, we aimed to give the next-generation probe a more druglike, nonpeptidic character to improve stability, bioavailability, and half-life in circulation. The Ellman laboratory recently used the substrate activity screening (SAS) approach to discover 1,4-disubstituted-1,2,3-triazole-based nonnatural aldehyde and nitrile inhibitors with high selectivity for cathepsin S over cathepsin B and L (Figure 1B) (Patterson et al., 2006; Wood et al., 2005). Triazoles, easily accessible through the Huisgen cycloaddition, have extensively been used as stable peptide bond mimics because of their similar electronic and structural characteristics. We therefore used these nonpeptidic triazole-based inhibitors as a starting point for design of our cathepsin S-directed ABPs (the synthesis of ABPs can be found in the [Supplemental Information, Figures S1–S3](#) available online). To determine the effect of the change of scaffold, we synthesized the fluorescent ABP BMV011 by replacing the norleucine P1 substituent by a lysine side chain to introduce the Cy5 fluorophore and the reversible electrophile by the irreversible 2,6-dimethyl benzoic AOMK (Figure 1C). We compared the labeling profile of BMV011 with that of the peptide-based probe GB123 (Blum et al., 2007) by exposing RAW cells (a mouse leukemic macrophage cell line) in culture to increasing concentrations of the probes. BMV011 and GB123 showed comparable labeling potency for cathepsin S ($EC_{50} = 1.0 \mu\text{M}$ and $= 1.3 \mu\text{M}$, respectively). However, the change in probe scaffold reduced labeling of cathepsin B ($EC_{50}(\text{BMV011}) = 6.0 \mu\text{M}$ and $EC_{50}(\text{GB123}) = 1.6 \mu\text{M}$) and cathepsin L (Figure 1D, top and middle panel), making BMV011 more selective for cathepsin S. Encouraged by these results, we synthesized the qABP BMV083, the nonpeptidic analog of GB137 (Figure 1C). The introduction of the quencher moiety further reduced the potency for cathepsin B, similar to the reduction of potency for cathepsin B seen for GB137 compared to GB123. This is probably due to steric hindrance with the occluding loop in the prime site pocket of cathepsin B. The primary target of BMV083 in RAW macrophages was cathepsin S, with a labeling potency similar to that of GB137 (Figure S4), and only minimal labeling of cathepsin L at high probe concentrations. Primary human monocyte-derived macrophages exposed to increasing concentrations of BMV083 showed a labeling profile similar to that observed in mouse macrophages, with cathepsin S being the primary target over cathepsins B and L (Figure 1E). Cathepsin S, in contrast to other cysteine cathepsin family members, has a wide pH optimum and is proteolytically active at both acidic and neutral pH (Kirschke et al., 1989). Consistent with this finding, we observed equal

labeling of cathepsin S by BMV083 in RAW cell lysate at acidic or neutral pH, and substantially diminished labeling of cathepsin B at neutral pH (Figure 1F).

Having determined the cell permeability and labeling specificity of BMV083, we wanted to visualize the dynamics of cysteine cathepsin labeling (unquenching) in living cells using confocal microscopy and correlate this signal with target labeling as assessed by SDS-PAGE analysis of the cell extracts. We exposed 4T1 mouse mammary adenocarcinoma cells to BMV083 and imaged at various time points (Figure 2A). Cy5 fluorescence signal increased over time and was primarily localized to a vesicular staining pattern. Fluorescent probe signals could be eliminated by pretreatment of the cells with a pan-cysteine cathepsin inhibitor (GB111-NH₂) (Blum et al., 2007), suggesting that probe activation was mediated by active cysteine cathepsin targets. Gel-based analysis of BMV083 labeling over the same time course showed that the time-dependent increase in Cy5 fluorescence activation correlated with labeling of cathepsin B and L. Furthermore, inhibition of target labeling by pretreatment with GB111-NH₂ also correlated with loss of imaging signals in the cells (Figure 2B). In agreement with its expression profile, no active cathepsin S was labeled in these cancer cells, which could explain the slow imaging and labeling kinetics.

To determine whether the probe could more effectively target cells with high cathepsin S levels, we next performed cell-based imaging studies in primary mouse bone marrow-derived macrophages (BMMs; Figures 2C and 2D). We found that in BMMs BMV083 is more rapidly unquenched compared to the 4T1 cells. Virtually all of the probe signal in the BMMs could be colocalized with a marker of lysosomal compartments (Figure 2D). On the basis of these findings, we wanted to see whether our cathepsin S-directed probe would be able to selectively label the cathepsin S-expressing cells in the presence of cells that predominantly express cathepsins B and L. Therefore, we cocultured BMMs with 4T1 cancer cells expressing both GFP and luciferase (4T1-luc-GFP) and imaged after probe treatment. Indeed, probe labeling was almost exclusively confined to the non-GFP-expressing BMMs (Figure 2E). Consistent with the imaging experiment, SDS-PAGE analysis of the labeled cysteine cathepsins in the coculture showed an intense fluorescent cathepsin S signal that was not present in the culture of the 4T1 cells alone (Figure 2B) and a labeling pattern very similar to that observed in the RAW mouse macrophage cell line (Figure 1D). Thus, the majority of labeling is derived from the macrophage population with only minimal labeling of cat B and L from the tumor cells.

In Vivo Properties of BMV083 in Tumor-Bearing Mice

Having determined the target preference and cell permeability of BMV083 *in vitro*, we next set out to establish the *in vivo* performance of the nonpeptidic qABP compared to the original peptidic GB137. We subcutaneously allografted athymic BALB/c nude mice with 4T1-luc-GFP cells. When tumors were established, we injected two cohorts of tumor-bearing mice via tail vein with equimolar amounts of either GB137 or BMV083 and noninvasively imaged the mice for Cy5 fluorescence over time (Figures 3A and 3B). Images of the whole mice are depicted in Figure S5A. As reported previously in xenograft tumor models, GB137 showed specific accumulation in tumors starting at early time points (Figure 3A) (Blum et al., 2007). The nonpeptidic

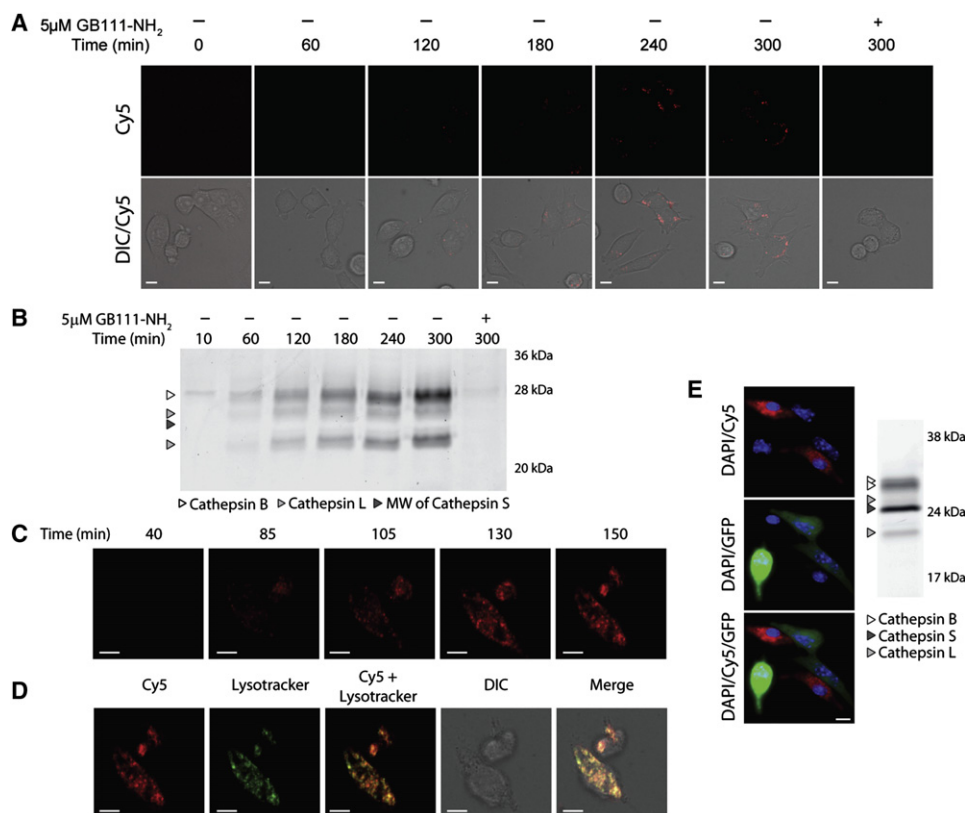


Figure 2. Live-Cell Imaging of Probe Labeling

(A–D) Live-cell confocal microscopy time course of BMV083 cysteine cathepsin labeling. (A) 4T1-luc-GFP cells were exposed to 1 μ M BMV083 and imaged over time. Where indicated, cells were preincubated with the cysteine cathepsin inhibitor GB111-NH₂ (5 μ M) for 1 hr before addition of probe (scale bar, 10 μ m). (B) 4T1-luc-GFP cells were exposed to 1 μ M BMV083 and harvested at indicated time points. Total protein (40 μ g) was resolved on 15% SDS-PAGE, and fluorescently labeled proteins were visualized by in-gel fluorescence scanning (MW = molecular weight). (C) Mouse primary bone marrow-derived macrophages were exposed to 1 μ M BMV083 and imaged over time (scale bar, 10 μ m). (D) After final time point, lysotracker was added (scale bar, 10 μ m). (E) 4T1-luc-GFP cells and mouse primary bone marrow-derived macrophages (2 to 1 ratio) were cocultured. After 90 hr, the cells were exposed to 1 μ M BMV083 for 2 hr. Cells were imaged (red, Cy5 fluorescence; green, GFP; blue, DAPI staining; scale bar, 10 μ m) and harvested. Total protein (40 μ g) was resolved on 15% SDS-PAGE, and fluorescently labeled proteins were visualized by in-gel fluorescence scanning.

counterpart BMV083, however, showed increased tumor-specific activation of fluorescence, resulting in substantially better contrast at all time points compared to GB137 (Figures 3A and 3B). The tumor-specific fluorescence intensity for BMV083, like GB137, peaked around the 4 hr time point (Figure 3C). Interestingly, unlike the GB137 tumor-specific fluorescence, which decreased over time, the BMV083 signal in tumors remained constant until the end of the time course. This could be due to enhanced stability of the free probe in vivo, resulting in extended labeling of newly synthesized pools of cysteine cathepsin. After the completion of the noninvasive optical imaging time course, we excised tumors and performed ex vivo imaging followed by homogenization and analysis of the fluorescently labeled protein content by SDS-PAGE (Figure 3D). Here, too, BMV083 had significantly increased overall cysteine cathepsin labeling compared to GB137. The in vivo targets for GB137 were primarily cathepsin B (38.0% \pm 5.7% [n = 5]) and L (33.6% \pm 2.3% [n = 5]) and, to a lesser extent, cathepsin S (28.4% \pm 1.8% [n = 5]). Although in cultured RAW cells, BMV083 almost exclusively labels cathepsin S in the presence of the more abundant cathepsin B (Figure 1D), in vivo

the primary target for BMV083 was cathepsin S (61.4% \pm 5.8% [n = 5]), with cathepsin B being the secondary target (38.6% \pm 5.2% [n = 5]). To determine the overall activity levels of the cysteine cathepsins in the 4T1 allografted tumors, we labeled tumor homogenates with the BODIPY TMR-labeled general cysteine cathepsin probe BMV099 (Figure 3D; for structure and characterization of BMV099, see Figure S6). Interestingly, this analysis indicated that the most active cysteine cathepsin in this tumor model is cathepsin B. Therefore, the relatively higher level of cathepsin B labeling by BMV083 in this model is likely because this is the most active cathepsin found in the tumor tissues. This result also stresses the fact that the selectivity of any inhibitor or substrate-based reporter highly depends on the experimental setup and the relative activity levels of other related (off-)targets in the tissue, cell, or sample being analyzed. The advantage of ABPs over substrate-based reporters is that the covalent irreversible nature of the labeling allows for biochemical validation and characterization of the in vivo observed probe fluorescence. Therefore, we could correlate the enhanced labeling signals by BMV083 in the live animals with whole-tissue fluorescence and relative levels of cysteine

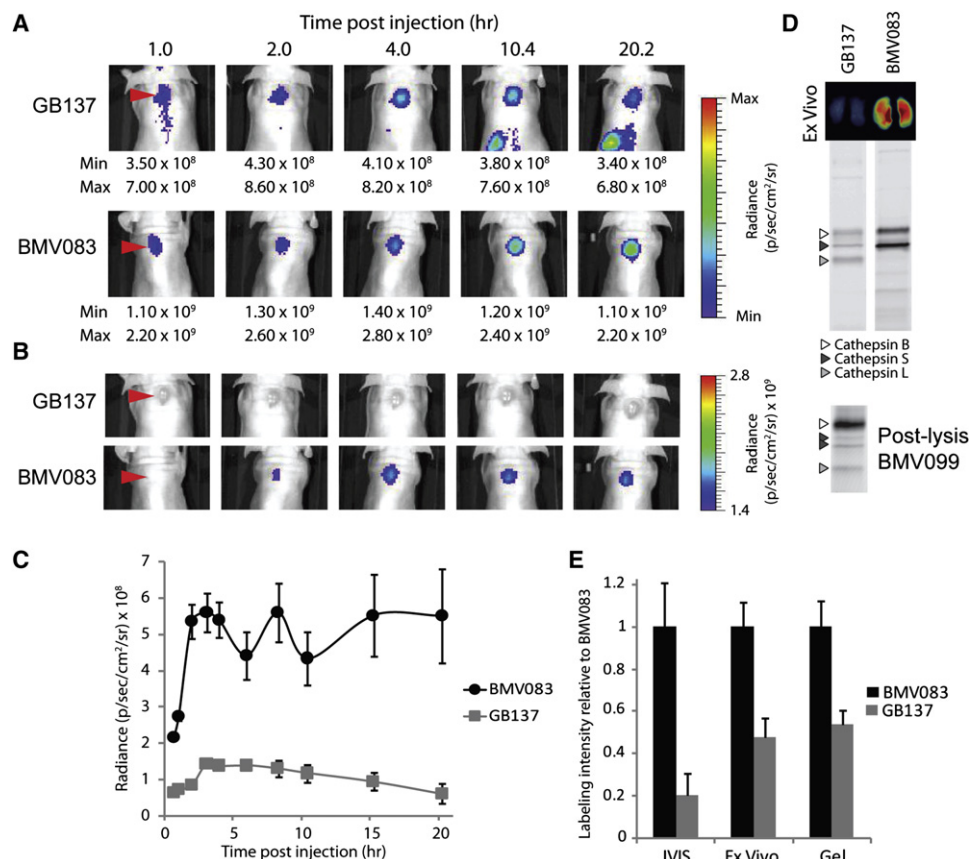


Figure 3. Noninvasive Optical Imaging of Cysteine Cathepsin Activity with Quenched Activity-Based Probes in Tumor-Bearing Mice

(A–C) 4T1-luc-GFP cells were injected subcutaneously on the back of athymic BALB/c nude mice, 8 days before imaging. GB137 or BMV083 (20 nmol) was injected (intravenous via tail vein), and fluorescent images of the living mice were taken over time. Red arrowheads indicate the tumor. (A) Optimal fluorescence contrast for individual time points. Maximum and minimum of scale bar depicted below image. (B) Comparison of GB137 and BMV083 fluorescence at fixed intensity. (C) Time-dependent tumor-specific fluorescence (tumor – background) for BMV083- and GB137-treated mice ($n = 5$; data represent mean values \pm standard errors).

(D) Ex vivo fluorescence imaging of cross-sections of excised tumors (upper panel). Tumor tissue was homogenized, and 40 μ g of total protein was resolved on 15% SDS-PAGE. In vivo fluorescently labeled proteins were visualized by in-gel fluorescence scanning (middle panel). 4T1 tumors were homogenized, and the active cysteine cathepsins were labeled with BMV099 (1 μ M for 1 hr; 80 μ g total protein) before the proteins were resolved on 15% SDS-PAGE. Fluorescently labeled proteins were visualized by in-gel fluorescence scanning (lower panel).

(E) Fluorescence intensity of end point of noninvasive optical imaging (shown in A), ex vivo tumor imaging, and in-gel fluorescence labeling (shown in D). Intensity relative to BMV083 is depicted ($n = 5$; data represent mean values \pm standard errors).

See also Figures S5 and S6.

cathepsin labeling observed for each probe by SDS-PAGE (Figure 3E).

The fact that GB137 and BMV083 have similar labeling potencies in vitro, yet the nonpeptidic probe was a more effective label in vivo, indicates that the nonpeptidic BMV083 has better in vivo properties compared to the peptidic GB137. Furthermore, liver and kidney fluorescence and cysteine cathepsin labeling were very similar for both probes (Figure S5B), indicating that BMV083 might circulate longer before being scavenged by these organs for clearance.

We next tested BMV083 in a more physiologically relevant orthotopic mouse breast cancer model. For this model, we implanted the 4T1 cell line isolated from a spontaneously arising BALB/c mammary tumor into the mammary fat pad of BALB/c mice with an intact immune system (Tao et al., 2008). This is especially important because host-tumor interactions and

immune cell infiltration in the tumor microenvironment are important factors in tumor development. We implanted 4T1-luc-GFP cells in the number 2 and 7 mammary fat pads, and tumor growth was monitored. We then intraperitoneally injected tumor-bearing mice with luciferin 10 hr after probe injection followed by imaging for bioluminescence and probe-derived Cy5 fluorescence (Figure 4A). The cancer cell-derived luciferase bioluminescence demarcated the tumor boundaries, and BMV083 activation was localized to the same area, indicating elevated cysteine cathepsin activity in the 4T1 tumor microenvironment. Interestingly, compared to the allograft model, fluorescence analysis of the SDS-PAGE-resolved protein content revealed cathepsin S as the major in vivo target for BMV083, with only minor labeling of cathepsin B (Figure 4B; cathepsin S represents $82.0\% \pm 0.7\%$ [$n = 6$] of total labeling intensity). Ex vivo labeling of the active cysteine cathepsin content in whole tumor homogenates with

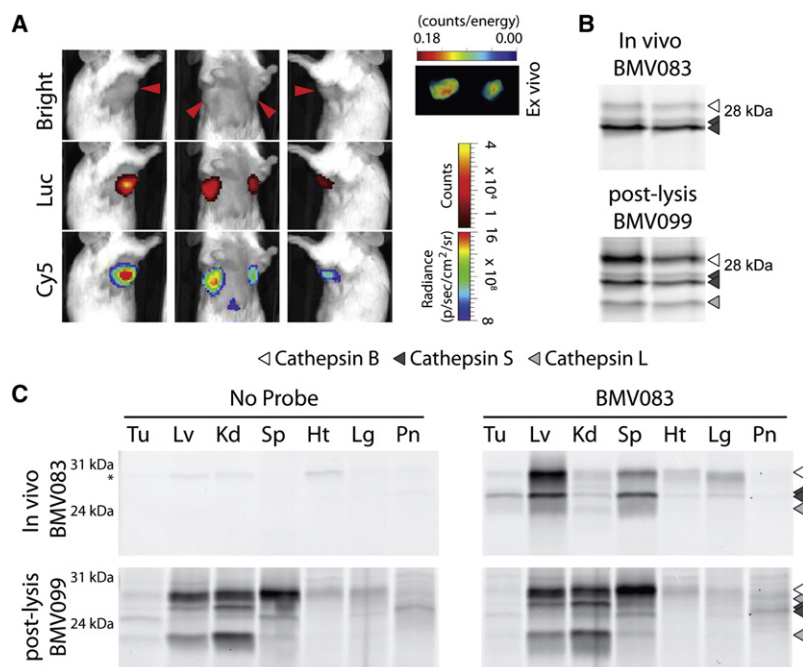


Figure 4. Noninvasive Optical Imaging of Cysteine Cathepsin Activity in a Syngeneic Orthotopic Model for Breast Cancer

(A) 4T1-luc-GFP cells were implanted into mammary fat pad numbers 2 and 7 of female BALB/c mice. On day 15, BMV083 (20 nmol) was injected (intravenous via tail vein). Ten hours after injection, the mice were injected with luciferin (intraperitoneal), and fluorescence and luminescence images of the living mice were taken (top panel, bright field image). Red arrowheads indicate the tumors; middle panel, luciferase bioluminescence (Luc); lower panel, BMV083 signal (Cy5). Ex vivo fluorescence image of excised tumors is shown.

(B) BMV083 labeling in tumors. The tumors were homogenized, and the active cysteine cathepsins were labeled with BMV099 (1 μ M for 1 hr; 40 μ g total protein) before the proteins were resolved on 15% SDS-PAGE. Fluorescently labeled proteins were visualized by in-gel fluorescence scanning (top panel, in vivo BMV083 labeling; bottom panel, ex vivo BMV099 labeling).

(C) In vivo BMV083 distribution. Organs were homogenized, and the active cysteine cathepsins were labeled with BMV099 (1 μ M for 1 hr; 40 μ g total protein) before the proteins were resolved on 15% SDS-PAGE. Fluorescently labeled proteins were visualized by in-gel fluorescence scanning. Top: in vivo BMV083 labeling. Bottom: ex vivo BMV099 labeling; *, autofluorescent band; Tu, tumor; Lv, liver; Kd, kidney; Sp, spleen; Ht, heart; Lg, lungs; Pn, pancreas.

See also Figure S6.

BMV099 indicated higher cathepsin S levels relative to cathepsin B compared to the xenograft model (Figure 3D), which likely explains why BMV083 labeled primarily cathepsin S in these tumors (Figure 4B). We performed a biodistribution study showing that, similar to what has been reported for GB137, the primary target organs of BMV083 are the liver and spleen (Figure 4C, upper panel). Ex vivo labeling of the active cysteine cathepsin content in whole-organ homogenates with BMV099 indicated a high degree of cysteine cathepsin activity in the organs that are strongly labeled in vivo (Figure 4C). Interestingly, although there are high levels of cysteine cathepsin activity in the kidney, it is not effectively targeted by BMV083, indicating specific partitioning of the probe to certain organs. On a similar note, the total cysteine cathepsin activity in the tumor was relatively low; however, the BMV083 signal in tumors is relatively high, again suggesting partitioning of the probe to specific tissues.

Cellular Source of BMV083 Labeling in Tumor-Bearing Mice

Fluorescent ABPs have previously been used to identify the cell populations within the tumor microenvironment that possess high levels of cysteine cathepsin activity. For example, fluorescence microscopy and immunofluorescence identified macrophages (F4/80⁺ cells) as the source of cysteine cathepsin activity in the RIP1-Tag2 mouse model of pancreatic islet cell cancer and in the MMTV-PyMT mouse model of mammary tumorigenesis treated with a nonquenched Cy3B labeled-DCG04 analog, "Cath-ABP" (Gocheva et al., 2010). In the same study, tumor-derived single-cell suspensions were labeled with the fluorescent Cath-ABP ex vivo and were analyzed by FACS. Again, macrophages were identified as the primary source of cysteine

cathepsin activity, although a nonmacrophage population was also found to take up probe. The authors suggested that this population might include cancer cells. Macrophage activation is a plastic and dynamic process that may change during processing of the tissue to generate single-cell suspensions for labeling and FACS analysis. Labeling of the tumor-derived cells in vivo is likely to be more physiologically relevant because the cysteine cathepsin targets are labeled in their native activation state. We therefore set out to determine the cellular source of BMV083-derived fluorescence in vivo. Because macrophages have previously been shown to possess high cysteine cathepsin activity in the tumor and because the amount of F4/80 expression is indicative of the macrophage activation state (i.e., M2 type macrophages have high F4/80 expression levels compared to a lower F4/80 expression on M1 type macrophages) (Fujiu et al., 2011), we analyzed four populations of cells from the 4T1-luc-GFP syngeneic orthotopic tumors for BMV083 fluorescence. These included two populations of F4/80-expressing cells (F4/80^{DIM} and F4/80^{HL}); cells that did not express GFP (F4/80⁻GFP⁻) and a population containing the GFP-expressing tumor cells (F4/80⁻GFP⁺) (Figure 5A). The F4/80^{HL} population was responsible for the majority of BMV083 activation, with a more than 12-fold increase in Cy5 mean fluorescence compared to vehicle treatment (Figure 5C, gray bars). To confirm this observation, we isolated the four cell populations in high purity using flow cytometric cell sorting, lysed equal cell numbers per population, and labeled the active cysteine cathepsins ex vivo with BMV099, followed by gel analysis of in vivo and ex vivo fluorescently labeled cysteine cathepsins (Figures 5B and 5C). This ex vivo labeling confirmed that in vivo probe labeling correlated with the absolute levels of cysteine cathepsins that were expressed and active in each cell population and was not the result

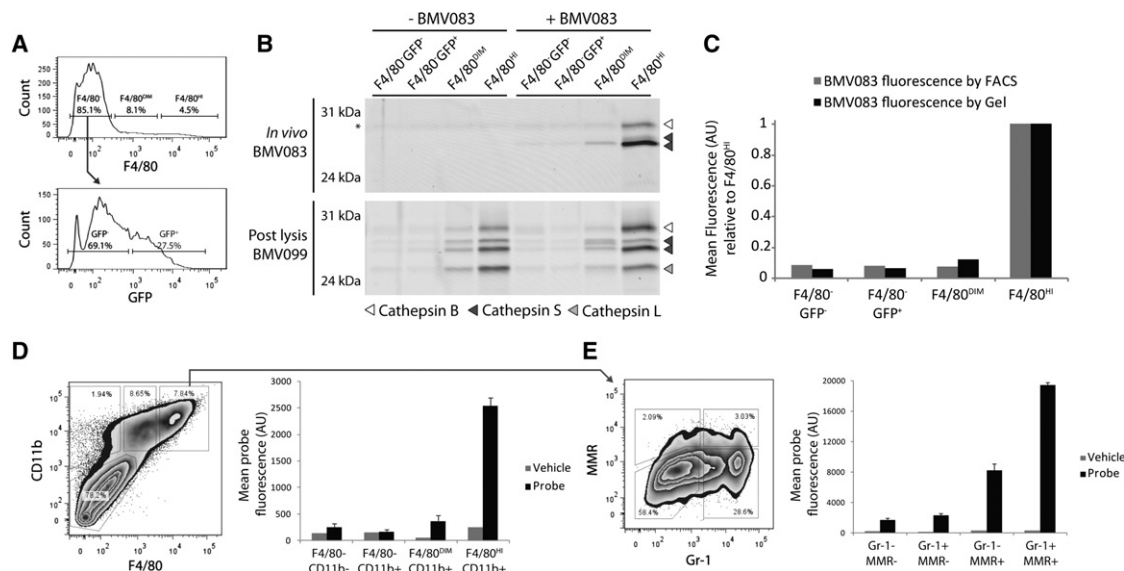


Figure 5. FACS Analysis to Determine Cellular Source of Cysteine Cathepsin Activity in 4T1 Syngeneic Breast Tumors

4T1 cells were implanted into mammary fat pad numbers 2 and 7 of female BALB/c mice. On day 15, BMV083 (20 nmol) was injected (intravenous via tail vein). Ten hours after injection, tumors were excised.

(A–C) Single-cell suspensions of excised tumors were analyzed by FACS for F4/80 and GFP expression and probe fluorescence. Four populations (F4/80⁺GFP⁻, F4/80⁺GFP⁺, F4/80⁺GFP⁺, and F4/80⁺) were analyzed and sorted. (A) F4/80 histogram (top panel) and GFP histogram of F4/80⁺ population (lower panel). (B) Equal cell numbers per population (depicted in A) of vehicle- and BMV083-treated mice were lysed, and the active cysteine cathepsins were labeled with BMV099 (1 μ M for 1 hr) before the proteins were resolved on 15% SDS-PAGE. Fluorescently labeled proteins were visualized by in-gel fluorescence scanning (top panel, in vivo BMV083 labeling; lower panel, ex vivo BMV099 labeling; *autofluorescent band). (C) Bar graph of quantification of mean Cy5 fluorescence per population (depicted in A) and in-gel fluorescence intensity of fluorescently labeled cysteine cathepsins (depicted in B) of vehicle- and BMV083-treated mice.

(D and E) Single-cell suspensions of excised tumors were analyzed by FACS for CD11b, F4/80, MMR, and Gr-1 expression and probe fluorescence. (D) FACS plot and bar graph of mean Cy5 fluorescence of four populations (CD11b⁺F4/80⁺, CD11b⁺F4/80⁺, CD11b⁺F4/80⁺, and CD11b⁺F4/80⁺) of vehicle- and BMV083-treated mice (n = 3; data represent mean values \pm standard errors). (E) FACS plot and bar graph of mean Cy5 fluorescence of four populations (MMR⁺Gr-1⁺, MMR⁺Gr-1⁺, MMR⁺Gr-1⁺, and MMR⁺Gr-1⁺) of CD11b⁺F4/80⁺ population depicted in (D) (n = 3; data represent mean values \pm standard errors).

See also Figure S6.

of unequal uptake of the probe by the different cell populations. Consistent with the overall mean fluorescence levels, the F4/80⁺ population showed the most robust labeling of cathepsin S and, to a lesser extent, cathepsin B, similar to the labeling profile observed in whole tumor homogenates (Figure 4B). We observed weaker labeling intensities in the other three populations, with a slightly more pronounced labeling in the F4/80⁺ population. We also observed a similar trend when analyzing the same gel for ex vivo BMV099-labeled cysteine cathepsins, indicating that the highest cysteine cathepsin activity was in the F4/80⁺ population. Interestingly, the degree of F4/80 expression seems to correlate with cysteine cathepsin activity, which suggests that cysteine cathepsin activity could be indicative of differences in activation phenotypes of myeloid-derived cells.

To further investigate which subpopulations of macrophages have the highest levels of cysteine cathepsin activity in the tumor microenvironment, we included three markers to identify the activation state of the labeled myeloid-derived cells. CD11b⁺Gr-1⁺ myeloid-derived suppressor cells (MDSCs) have been shown to possess high cysteine cathepsin activity in the APC^{Δ468} mouse model of hereditary polyposis (Gounaris et al., 2008). It is also generally believed that tumor-associated macrophages (TAMs) possess an M2-like macrophage character (Mukhtar et al., 2011). We therefore used the macrophage

mannose receptor (MMR, also known as CD206), which is elevated in M2 compared to M1 macrophages, as a marker for the different macrophage populations (Mukhtar et al., 2011). Because of recent reports concerning the immunogenicity of GFP in BALB/c mice (Bosiljic et al., 2011), we also implanted 4T1 cells lacking the GFP reporter in the number 2 and 7 mammary fat pads of BALB/c mice. We systemically administered BMV083 via tail vein injection and excised tumors at 10 hr after injection to generate single-cell suspensions. Consistent with the results in the 4T1-luc-GFP tumors, we observed the highest levels of probe activation in the F4/80⁺ cell population, as determined by FACS analysis (CD11b⁺F4/80⁺; Figure 5D). To further characterize the probe-positive cells, we analyzed the CD11b⁺F4/80⁺ population for Gr-1 and MMR expression (Figure 5E). This analysis determined that BMV083 labeling occurs most predominantly in CD11b⁺F4/80⁺ cells that express MMR. Furthermore, of the CD11b⁺F4/80⁺MMR⁺ cells, the Gr-1-expressing population showed more than 2-fold higher probe accumulation compared to the Gr-1-negative population. These data indicate that the CD11b⁺F4/80⁺MMR⁺Gr-1⁺ cells are the major cellular source of cathepsin S activity in the 4T1 tumor microenvironment. CD11b⁺F4/80⁺Gr-1⁺ cells in mouse models of cancer have been defined as myeloid-derived suppressor cells (MDSCs). In response to TLR4 ligands released from damaged tumor cells, MDSCs

inhibit antitumor T cell responses by inducing apoptosis in activated T cells (Liu et al., 2010).

The data presented here show that the cathepsin S-directed nonpeptidic NIRF qABP BMV083 has improved in vivo properties compared to previously developed ABPs. In the syngeneic 4T1 mouse model for breast cancer, this probe primarily reports on the cysteine cathepsin activity in the tumor-promoting M2 type CD11b⁺F4/80⁺MMR⁺ macrophages. Of these, the Gr-1 expressing MDSCs constitute the highest degree of probe activation. Because infiltration of these cell types into the tumor microenvironment plays an important role in tumor development and because their occurrence is correlated with poor prognosis and negative effects on tumor immunotherapy, BMV083 could be of prognostic value. Furthermore, BMV083 may find application in the study of the role of cathepsin S in other diseases, such as atherosclerosis, inflammatory disorders, and rheumatoid arthritis (Conus and Simon, 2010; Quillard et al., 2011; Reiser et al., 2010).

SIGNIFICANCE

Cells of myeloid origin are major contributors to the development of many disease states. In vivo imaging tools for macrophages could greatly facilitate the study of their role in disease pathology (Quillard et al., 2011). The consequence of macrophage infiltration in tumor development has been extensively studied in the last few decades. Some consider cancer-related inflammation the seventh hallmark of cancer (Colotta et al., 2009), whereas others consider tumor-promoting inflammation a hallmark enabling characteristic (Hanahan and Weinberg, 2011). Regardless, it is clear that tumor-associated macrophages play important roles in tumor development and have been shown to promote angiogenesis, tumor growth, and invasiveness (Mantovani et al., 2008; Qian and Pollard, 2010). Furthermore, macrophage infiltration into tumors has been correlated with poor clinical outcome in several cancers (Bingle et al., 2002; Kurahara et al., 2011; Steidl et al., 2010). The activation state acquired by the infiltrating macrophage in the tumor microenvironment determines its effect on the tumor. Macrophages with the M2 phenotype exhibit tumor-promoting characteristics, whereas M1 macrophages are able to eradicate tumors. The macrophage activation state is plastic, and tumor-promoting macrophages can be converted into tumor-killing M1 macrophages, for example, by local treatment with granulocyte macrophage colony-stimulating factor (Eubank et al., 2009). Noninvasive imaging tools that report on the tumor-promoting M2 type tumor-associated macrophage content could therefore be of prognostic value. This manuscript presents the synthesis and characterization of BMV083, a nonpeptidic cathepsin S-directed NIRF qABP. This probe shows improved in vivo properties compared to our previous generation of peptidic quenched probes, with higher tumor-specific fluorescence in noninvasive optical imaging experiments. Furthermore, the primary cellular source of probe labeling in the tumor was determined to be macrophages with an M2 phenotype, making BMV083 a potential in vivo reporter for tumor-associated macrophages that primarily promote tumor growth.

EXPERIMENTAL PROCEDURES

General

All resins and reagents were purchased from commercial suppliers and used without further purifications. All solvents used were HPLC grade. All water-sensitive reactions were performed in anhydrous solvents under positive pressure of argon. Reactions were analyzed by LC-MS using an API 150EX single-quadrupole mass spectrometer (Applied Biosystems). Reverse-phase HPLC was conducted with an ÄKTA explorer 100 (Amersham Pharmacia Biotech) using C18 columns. NMR spectra were recorded on a Varian 400 MHz (400/100) or a Varian Inova 600 MHz (600/125 MHz) equipped with a pulsed field gradient accessory. Chemical shifts are given in ppm (δ) relative to tetramethylsilane as an internal standard. Coupling constants are given in Hz. Fluorescent gels were scanned using a Typhoon 9400 flatbed laser scanner (GE Healthcare). Statistical analysis was performed using Microsoft Excel, and s.e.m. was calculated by dividing the s.d. by the square root of *n*. Fluorescent images were acquired on a Zeiss Axiovert 200 M inverted microscope equipped with a 10 \times , 40 \times , and 63 \times objective (Carl Zeiss). Slidebook software was used to control the microscope and camera and for data analysis (Intelligent Imaging Innovations).

ABP Synthesis

For the synthesis and characterization of BMV011, BMV083, and BMV099 see the Supplemental Information.

Cell Culture and Labeling of Living Cells

RAW cells were cultured in DMEM (GIBCO) supplemented with 10% fetal bovine serum (FBS; GIBCO), 100 U/ml penicillin, and 100 μ g/ml streptomycin (GIBCO). 4T1 cells (ATCC) and 4T1-luc-GFP cells were cultured in RPMI (GIBCO) supplemented with 10% FBS (GIBCO), 100 U/ml penicillin, and 100 μ g/ml streptomycin (GIBCO). Primary mouse bone marrow-derived macrophages (BMDMs) were isolated and cultured as previously described (Broz et al., 2010). Briefly, bone marrow was isolated from femurs of 6–8-week-old C57BL/6 mice (The Jackson Laboratory, Bar Harbor, ME). BMDMs were differentiated in DMEM (GIBCO) with 10% FBS and 10% MCSF-containing cell supernatant for 6 days. The cells were then frozen in FBS with 10% DMSO and thawed prior to use. All cells were cultured in a 5% CO₂ humidified incubator at 37°C. Cocultures of 4T1-luc-GFP cells and BMDMs were established by seeding cells in a 2 to 1 ratio and culture in DMEM supplemented 10% heat-inactivated FBS (GIBCO), 100 U/ml penicillin, and 100 μ g/ml streptomycin (GIBCO) for 90 hr. For the derivation of primary human macrophages, leukocyte reduction system chambers were obtained from anonymous blood donations at the Stanford Blood Center. Peripheral blood mononuclear cells were isolated by Ficoll gradient centrifugation, and then monocytes were purified on an AutoMACS (Miltenyi) by positive selection using CD14 microbeads (Miltenyi). Monocytes were differentiated to macrophages by culturing for 9 days in Iscove's Modified Dulbecco's Medium supplemented with 10% AB human serum (Invitrogen), 100 U/ml penicillin, and 100 μ g/ml streptomycin (Invitrogen). For intact cell labeling, cells were exposed to probe or inhibitor (1 hr preincubation) where indicated (1000 \times in DMSO) for the indicated times. The cells were harvested, washed with PBS, resuspended in hypotonic lysis buffer (50 mM PIPES [pH 7.4], 10 mM KCl, 5 mM MgCl₂, 2 mM EDTA, 4 mM DTT, and 1% NP-40), put on ice for 15 min, and centrifuged at 4°C, and supernatants were collected. Total protein (40 μ g) was denatured with SDS-sample buffer and resolved by SDS-PAGE (15%), and labeled proteases were visualized by scanning the gel with a Typhoon imager (GE Healthcare). Labeling intensities were quantified using Image J software. For lysate labeling of active cysteine cathepsins, cells were lysed in citrate buffer (50 mM citrate [pH 5.5 or pH 7.0], 5 mM DDT, 0.5% CHAPS, and 0.1% Triton X). Total protein (35 or 40 μ g) was labeled with the indicated probe at indicated concentration for 1 hr at 37°C before analysis as described above. For live cell confocal microscopy, cells were seeded in an 8 well Lab-Tek chambered coverglass system.

Animal Models

All animal care and experimentation was conducted in accord with current National Institutes of Health and Stanford University Institutional Animal

Care and Use Committee guidelines. For the 4T1 allograft model, female athymic nude mice (nu/nu) (6–8 weeks old, Charles River Laboratories) were injected subcutaneously with 3×10^5 4T1-luc-GFP cells in PBS under isoflurane anesthesia, and tumor growth was monitored. On day 8, GB137 or BMV083 (20 nmol; 0.8 nmol g^{-1}) was administered via tail vein in 100 μL volume (60% DMSO in PBS). After injection, mice were imaged noninvasively at indicated time points using an IVIS 100 system. After the last time point, the mice were anesthetized with isoflurane and killed by cervical dislocation. Tumors were removed and imaged ex vivo using an FMT 2500 (Perkin Elmer), and the tissue was homogenized in muscle lysis buffer (1% Triton X-100, 0.1% SDS, and 0.5% sodium deoxycholate in PBS [pH 7.4]). Proteins (40 μg) were resolved by SDS-PAGE (15%), and labeled proteases were visualized by scanning the gel with a Typhoon imager (GE Healthcare). Labeling intensities were quantified using Image J software. For the orthotopic mouse breast cancer model, female BALB/c mice (6–8 weeks old, The Jackson Laboratory) were injected in the fat pad numbers 2 and 7 with 1×10^5 4T1-luc-GFP or 4T1 cells (ATCC) in PBS, and tumor growth was monitored. On day 15, the mice were injected with BMV083 and analyzed as described above. For ex vivo labeling of active cysteine cathepsins, tissues were removed and homogenized in citrate buffer (50 mM Citrate [pH 5.5], 5 mM DDT, 0.5% CHAPS, and 0.1% Triton X). Total protein (40 μg) was labeled with BMV099 (1 μM final concentration) for 1 hr at 37°C before analysis as described above.

Antibody Staining and Flow Cytometry

The tumor tissue was dissected from the mouse, sliced into 1–2 mm³ pieces with a razor blade, and digested at 37°C for approximately 2 hr, with frequent pipetting in Advanced DMEM/F12 (Invitrogen) with 2 mM Glutamax (Invitrogen), 120 $\mu\text{g}/\text{mL}$ penicillin, 100 $\mu\text{g}/\text{mL}$ streptomycin, and 0.25 $\mu\text{g}/\text{mL}$ amphotericin-B (PSA), with 200 U/mL Collagenase type III (Worthington, Lakewood, NJ) and 100 U/mL DNase I (Worthington). After the digestion reached completion, cells were filtered with 40 μm nylon mesh (BD Biosciences), washed, and resuspended at a density of 10×10^6 cells/mL in cold HBBS (Invitrogen) with PSA and 2% heat inactivated fetal calf serum. To reduce nonspecific binding, cells suspended in staining buffer were blocked on ice for 10 min with 10 mg/mL rat IgG (Sigma) at 1:1000. Cells were then stained, in the dark, on ice for 30 min with optimal antibodies concentrations, which was determined by titration experiments. Antibodies used include PE-Cy7 anti-mouse F4/80 (BM8), PE-Cy5 anti-mouse CD11b (M1/70) (ebioscience), FITC anti-mouse MMR (C068C2), and PE anti-mouse Gr-1 (RB6-8C5) (BioLegend). Flow cytometry was performed with a BD FACS Aria II with FACS Diva software. Data analysis was performed using FlowJo. For all experiments, side scatter and forward scatter profiles (area and width) were used to eliminate debris and cell doublets. Dead cells were eliminated by excluding DAPI⁺ cells (Molecular Probes).

SUPPLEMENTAL INFORMATION

Supplemental Information includes six figures and can be found with this article online at doi:10.1016/j.chembiol.2012.03.012.

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